



STIC Search Report

Biotech-Chem Library

STIC Database Tracking Number: 151270

TO: Janet Epps-Ford
Location: rem/2c05/2c18
Art Unit: 1635

April 29, 2005

Case Serial Number: 10/086062

From: P. Sheppard
Location: Remsen Building
Phone: (571) 272-2529

sheppard@uspto.gov

Search Notes

04/29/05

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STIC-Biotech/ChemLib

151270

me

From: Epps-Ford, Janet
Sent: Wednesday, April 20, 2005 5:14 PM
To: STIC-Biotech/ChemLib
Subject: Sequence search.

Application 10/086,062

Please search SEQ ID NO: 4 in all pending and published nucleic acid databases.

Thanks,

Janet L. Epps-Ford, Ph.D.

Art Unit 1635

Mailbox: Remsen 2C18

Office: Remsen 2C05

Phone: 571-272-0757

Fax: 571-273-0757

STAFF USE ONLY

Searcher: _____
Searcher Phone: 2-_____
Date Searcher Picked up: _____
Date Completed: _____
Searcher Prep/Rev. Time: _____
Online Time: _____

Type of Search

NA#: _____ AA#: _____
Interference: _____ SPDI: _____
S/L: _____ Oligomer: _____
Encode/Transl: _____
Structure#: _____ Text: _____
Inventor: _____ Litigation: _____

Vendors and cost where applicable

STN: _____
DIALOG: _____
QUESTEL/ORBIT: _____
LEXIS/NEXIS: _____
SEQUENCE SYSTEM: _____
WWW/Internet: _____
Other(Specify): _____

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REFERENCE	Mammalia; Eutheria; Primates; Carnivora; Hominiidae; Homo.
AUTHORS	Moensner,J., Tan,F., Marra,M., Kucaba,T., Yandell,M., Martin,J., Meth,G., Bowles,L., Wyle,T., Bowers,Y., Steptoe,M., Treising,B., Gibbons,R., Allen,M., Underwood,K., Chappell,J., Person,B., Gibbons,M., Harvey,N., Pape,D., Chamberlain,A., Morales,R., Schurk,R., Ritter,E., Kohn,S., Swaller,T., Behrmer,K., Hillier,L., Wilson,R. and Waterston,R.
TITLE	Full Clone Sequencing of the Longest Available Member from Each Unigene Cluster
JOURNAL	Submitted (24-AUG-1998) Department of Genetics, Washington University, 444 Forest Park Avenue, St. Louis, Missouri 63108, USA
REFERENCE	Submitted By: Genome Sequencing Center Department of Genetics Washington University St. Louis MO 63108, USA http://genome.wustl.edu/gsc mailto:est@watson.wustl.edu
AUTHORS	NOTICE: This sequence represents the full insert of this cDNA. No attempt has been made to verify whether this corresponds to the full-length of the original mRNA from which it was derived. We have tried to obtain double-stranded, or double chemistry sequence across the entire clone, but potentially, there are areas in the sequence where this level of coverage was not achieved. Nevertheless, we are confident of the accuracy of this sequence as all regions of low quality, as defined by PHRAP (P. Green, in preparation), were visually inspected and edited accordingly. The consensus quality values for this sequence have been submitted separately.
COMMENT	The location of this clone is unknown.
FEATURES	Location/Qualifiers 1..595 /organism="Homo sapiens" /mol_type="mRNA" /db_xref="taxon:9606" /clone="IMAGE:201721" /clone_lib="Scarses_fetal_liver_spleen_mRFLS" 9..338 /rnc_family="L2"
SOURCE	Query Match 68.0%; Pred. 20.4; DB 9; Length 595; Best Local Similarity 80.0%; Score.No.2.9e+02; Matches 24; Conservative 0; Mismatches 6; Indels 0; Gaps 0;
ORIGIN	repeat_region CTGACCCCTCTCGACTCGAGATGCCGCT 30 566 CTGACCCCTCTCGACTCGAGATGCCGCT 537
OY	AL136103 98348 bp DNA linear PRI 04-JAN-2001
LOCUS	Human DNA sequence from clone RPL-250B11 on chromosome 10 contains STS and GSSs, complete sequence.
DEFINITION	AL136103
ACCESSION	AL136103.24 GI:9581541
VERSION	HTG.
KEYWORDS	Homo sapiens (human)
SOURCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Carnivora; Hominiidae; Homo.
ORGANISM	1 (bases 1 to 98348) Phillimore,B. Direct Submission Submitted (04-JAN-2001) Sanger Centre, Hinxton, Cambridgeshire,

COMMENT

CB30 1SA. UK. E-mail enquiries: humquerry@sanger.ac.uk
 requests: clonerquests@sanger.ac.uk
 On Jul 28, 2000 this sequence version replaced g1:9501151.
 During sequence assembly data is compared from overlapping clones.
 Where differences are found these are annotated as variations
 together with a note of the overlapping clone name. Note that the
 variation annotation may not be found in the sequence submission
 corresponding to the overlapping clone, as we submit sequences with
 only a small overlap as described above.

The following abbreviations are used to associate primary accession
 numbers given in the feature table with their source databases:
 Em, EMBL; Sw, SWISSPROT; Tr, TREMBL; Wp, WORMPEP. Information
 on the WORMPEP database can be found at
http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence is
 the entire insert of clone RP1-250B11 This sequence has been
 finished according to sequence map criteria as follows. An attempt
 is made to resolve all sequencing problems, such as compressions
 and repeats, but not necessarily within known annotated repeat
 sequence elements. Where the sequence is ambiguous, there is an
 annotation using the 'unsure' feature key. This sequence was
 generated from part of bacterial clone contigs of human chromosome
 10, constructed by the Sanger Centre Chromosome 10 Mapping Group.
 Further information can be found at
<http://www.sanger.ac.uk/HGP/Chr10>
 RP1-250B11 is from the library RPCI-1 constructed by the group of
 Pieter de Jong. For further details see
<http://www.chori.org/bacpac/home.htm>
 VECTOR: pCYPAC2.

FEATURES

source

1..98348
 /organism="Homo sapiens"
 /mol_type="genomic DNA"
 /db_xref="taxon:9606"
 /chromosome="10"
 /clone="RP1-250B11"
 /clone_1fb="RPCI-1"

1..34
 /note="Single clone region. Reads derived from clone PCR.
 Assembly consistent with restriction digest."

repeat_region
 2459..2765
 /note="AluX repeat: matches 1..308 of consensus"

repeat_region
 3087..3267
 /note="MER39b repeat: matches 355..546 of consensus"

repeat_region
 3269..3606
 /note="MER3 repeat: matches 13..380 of consensus"

repeat_region
 6106..6337
 /note="L2 repeat: matches 2435..2705 of consensus"

repeat_region
 6364..6662
 /note="AluSg repeat: matches 1..298 of consensus"

repeat_region
 complement(8333..8799)
 /note="match: GSS: Em:AQ0403443"

misc_feature
 complement(8375..8758)
 /note="match: GSS: Em:AQ264393"

misc_feature
 8802..9234
 /note="match: GSS: Em:AQ150662"

misc_feature
 9268..9356
 /note="match: STS: Em:G28019"

repeat_region
 11809..12092
 /note="AluX repeat: matches 1..284 of consensus"

repeat_region
 12205..12722
 /note="L1 repeat: matches 5285..5811 of consensus"

repeat_region
 14113..14171
 /note="L2 repeat: matches 2692..2750 of consensus"

repeat_region
 14548..14846
 /note="AluY repeat: matches 1..299 of consensus"

repeat_region
 15883..16169
 /note="AluX repeat: matches 1..287 of consensus"

repeat_region
 17146..18557
 /note="MER52A repeat: matches 1..1755 of consensus"

repeat_region
 18559..18713
 /note="MIR repeat: matches 6..160 of consensus"

repeat_region
 18955..19381
 /note="MER63 repeat: matches 590..1061 of consensus"

```

repeat_region 19384..19519
/note="FLAM C repeat: matches 3..137 of consensus"
repeat_region 19713..19828
/note="MER63 repeat: matches 463..577 of consensus"
repeat_region 19826..20117
/note="MER63 repeat: matches 2..307 of consensus"
repeat_region 20355..20391
/note="MIR repeat: matches 88..156 of consensus"
repeat_region 20440..20701
/note="MIR repeat: matches 2..248 of consensus"
misc_feature 22321..22631
/note="match: STS: Em:HSJ10C4"
misc_feature 25725..26188
/note="match: GSS: Em:AQ211002"
misc_feature 25738..25978
/note="match: GSS: Em:AQ823705"
misc_feature 26548..27067
/note="match: GSS: Em:AQ798605"
misc_feature 26564..27250
/note="match: GSS: Em:AQ055423"
misc_feature 27549..27913
/note="match: GSS: Em:B87793"
repeat_region 28724..28952
/note="L2 repeat: matches 2167..2416 of consensus"
misc_feature complement(29565..30037)
/note="match: GSS: Em:AQ155973"
misc_feature 30073..30670
/note="match: GSS: Em:AQ195052"
misc_feature complement(31752..32121)
/note="match: GSS: Em:AQ099305"
misc_feature complement(32344..32928)
/note="match: GSS: Em:AQ485533"
misc_feature 32930..33368
/note="match: GSS: Em:AQ424293"
repeat_region 34554..34583
/note="15 copies 2 mer ca 100% conserved"
repeat_region 35059..35135
/note="L2 repeat: matches 2417..2500 of consensus"
misc_feature complement(35681..36191)
/note="match: GSS: Em:B55923"
misc_feature complement(35777..36102)
/note="match: GSS: Em:AQ223744"
misc_feature complement(35861..36186)
/note="match: GSS: Em:AQ803876"
repeat_region 36692..36800
/note="MIR repeat: matches 13..123 of consensus"
repeat_region 40461..40765
/note="AluX repeat: matches 1..293 of consensus"
repeat_region 41312..41502
/note="MIR repeat: matches 8..192 of consensus"
repeat_region 41785..41840
/note="28 copies 2 mer tg 85% conserved"
repeat_region 43022..43334
/note="AluX repeat: matches 3..312 of consensus"
misc_feature complement(44477..45066)
/note="match: GSS: Em:AQ540223"
misc_feature complement(44671..45063)
/note="match: GSS: Em:AQ337658"
repeat_region 45077..45157
/note="MER57-internal repeat: matches 7151..7230 of consensus"
repeat_region 45158..45304
/note="LIP repeat: matches 5050..5197 of consensus"
repeat_region 45284..45621
/note="LIP repeat: matches 5476..5812 of consensus"
misc_feature complement(45941..46452)
/note="match: GSS: Em:AQ832143"
repeat_region 48564..48709
/note="L2 repeat: matches 2596..2749 of consensus"
repeat_region 48720..49082
/note="THE1B repeat: matches 1..364 of consensus"
repeat_region 49176..49629
/note="MLTID repeat: matches 1..502 of consensus"

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repeat_region 49784..49912
/note="FLAM A repeat: matches 8..132 of consensus"
repeat_region 49914..50164
/note="MIR repeat: matches 2..261 of consensus"
repeat_region 52850..52918
/note="L2 repeat: matches 2635..2706 of consensus"
misc_feature 53092..53613
/note="match: GSS: Em:AQ800606"
misc_feature 53110..53608
/note="match: GSS: Em:AQ172250"
misc_feature 53151..53604
/note="match: GSS: Em:AQ170181"
misc_feature 53165..53586
/note="match: GSS: Em:AQ545768"
misc_feature 53179..53355
/note="match: GSS: Em:AQ898990"
repeat_region 53378..53834
/note="match: GSS: Em:AQ898990"
repeat_region 53776..53958
/note="MIR repeat: matches 41..250 of consensus"
misc_feature complement(53900..54343)
/note="match: GSS: Em:AQ400632"
repeat_region 53959..54203
/note="AluX repeat: matches 65..308 of consensus"
repeat_region 54204..54216
/note="MIR repeat: matches 250..260 of consensus"
misc_feature 54361..54767
/note="match: GSS: Em:AQ729256"
repeat_region 54372..54565
/note="match: GSS: Em:AQ729256"
misc_feature 54440..54730
/note="match: GSS: Em:AQ084956"
misc_feature complement(55008..55535)
/note="match: GSS: Em:AQ556262"
misc_feature 55610..55935
/note="match: GSS: Em:AQ594863"
repeat_region 56871..57293
/note="LMD repeat: matches 883..1319 of consensus"
repeat_region 57297..57602
/note="AluX repeat: matches 1..305 of consensus"

Query Match 68.0%; Score 20.4; DB 9; Length 98348;
Best Local Similarity 80.0%; Pred. No. 2e+02;
Matches 24; Conservative 0; Mismatches 6; Indels 0; Gaps 0;

Qy 1 CTGAGCCCTCTGACCTGAGAGTCCGCT 30
Db 62965 CTGAGCCCTCTCGCTGAGACTTCTCT 62994

RESULT 4
AC012048 178965 bp DNA linear PRI 20-APR-2002
LOCUS Homo sapiens chromosome 10 clone RP11-43N22, complete sequence.
DEFINITION AC012048
ACCESSION AC012048.11 GI:19744964
VERSION 1
KEYWORDS HTG.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Mammalia; Eutheria; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
REFERENCE 1 (bases 1 to 178965)
AUTHORS Smith,D.R.
TITLE Genome Therapeutics Corporation Sequencing Center: Human Genome
Sequence Data
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 178965)
AUTHORS Smith,D.R.
TITLE Direct Submission
JOURNAL Submitted (19-OCT-1999) Genome Therapeutics Corporation, 100 Beaver
Street, Waltham, MA 02453, USA
REFERENCE 3 (bases 1 to 178965)
AUTHORS Smith,D.R.

```

TITLE Direct Submission
JOURNAL Submitted (16-AUG-2001) Genome Therapeutics Corporation, 100 Beaver Street, Waltham, MA 02453, USA
REFERENCE 4 (bases 1 to 178965)
AUTHORS Smith,D.R.
JOURNAL Direct Submission
SUBMITTED (27-MAR-2002) Genome Therapeutics Corporation, 100 Beaver Street, Waltham, MA 02453, USA
REFERENCE 5 (bases 1 to 178965)
AUTHORS Smith,D.R.
TITLE Direct Submission
JOURNAL Submitted (28-MAR-2002) Genome Therapeutics Corporation, 100 Beaver Street, Waltham, MA 02453, USA
REFERENCE 6 (bases 1 to 178965)
AUTHORS Smith,D.R.
TITLE Direct Submission
JOURNAL Submitted (20-APR-2002) Genome Therapeutics Corporation, 100 Beaver Street, Waltham, MA 02453, USA
COMMENT On Mar 27, 2002 this sequence version replaced g1:15193325.
FEATURES
source
1. 178965
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/chromosome="10"
/clone="RP11-43N22"
/clone_1b="RP11-11"

ORIGIN
Query Match 68.0%; Score 20.4; DB 9; Length 178965;
Best Local Similarity 80.0%; Pred. No. 1.9e+02;
Matches 24; Conservative 0; Mismatches 6; Indels 0; Gaps 0;
QY 1 CTGACCCCTCTCGACTCGAGTTCGCT 30
DB 32913 CTGAGCCCTCTCGCTGAGACTCTCT 32942

RESULT 5
AC099767/c 23211 bp DNA linear INV 20-NOV-2001
LOCUS Caenorhabditis briggsae cosmid G24F01, complete sequence.
DEFINITION AC099767
AC099767.1 GI:17017639
HTG.
Caenorhabditis briggsae
Caenorhabditis briggsae
Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida;
Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
1 (bases 1 to 23211)
Washington University Genome Sequencing Center.
The C. briggsae Genome Sequencing Project
Unpublished
2 (bases 1 to 23211)
Waterston,R.
Direct Submission
Unpublished
3 (bases 1 to 23211)
Waterston,R.
Direct Submission
Submitted (20-NOV-2001) Department of Genetics, Washington University, Genome Sequencing Center, 444 Forest Park Avenue, St. Louis, MO 63110, USA
Submitted by:
Genome Sequencing Center
Department of Genetics, Washington University
St. Louis, MO 63110, USA
email: rw@nematode.wustl.edu

COMMENT
NOTICE: This sequence may not be the entire insert of this clone. It may be shorter because we only sequence overlapping sections once, or longer because we provide a small overlap between neighboring submissions.

This sequence was finished as follows unless otherwise noted: all regions were double stranded, sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one m13 subclone.
Location/Qualifiers
1. 23211
/organism="Caenorhabditis briggsae"
/mol_type="genomic DNA"
/strain="GujAra1 G16"
/db_xref="taxon:6238"

ORIGIN
Query Match 67.3%; Score 20.2; DB 3; Length 23211;
Best Local Similarity 86.0%; Pred. No. 2.7e+02;
Matches 22; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 4 GACCCCTCTCGACTCGAGTTCG 28
DB 14462 GATTCCTCTCACTCGAGGTTCCG 14438

RESULT 6
AB038027/c 1503 bp DNA linear BCT 18-FEB-2000
LOCUS Vibrio sp. SG128 gene for 16S rRNA, partial sequence.
DEFINITION AB038027
AB038027.1 GI:7007473
VERSION 16S ribosomal RNA.
KEYWORDS Vibrio sp. SG128
SOURCE Vibrio sp. SG128
ORGANISM Bacteria; Proteobacteria; Gammaproteobacteria; Vibrionales; Vibrionaceae; Vibrio.
1 (bases 1 to 1503)
Urakawa,H.
16S rRNA gene of marine bacterium
Published Only in Database (2000)
2 (bases 1 to 1503)
Urakawa,H.
Direct Submission
Submitted (05-FEB-2000) Hidetoshi Urakawa, Northwestern University, Department of Civil Engineering, Technological Institute 2145 Sheridan Road, Evanston, Illinois 60208-3109, USA
(E-mail:h-urakawa@nwu.edu, Tel:+1-847-467-5710, Fax:+1-847-491-4011)
Location/Qualifiers
1. 1503
/organism="Vibrio sp. SG128"
/mol_type="genomic DNA"
/strain="SG128"
/db_xref="taxon:115126"
/product="16S ribosomal RNA"

FEATURES
source
1. 1503
/organism="Vibrio sp. SG128"
/mol_type="genomic DNA"
/strain="SG128"
/db_xref="taxon:115126"
/product="16S ribosomal RNA"

ORIGIN
Query Match 66.7%; Score 20; DB 1; Length 1503;
Best Local Similarity 82.1%; Pred. No. 4.1e+02;
Matches 23; Conservative 0; Mismatches 5; Indels 0; Gaps 0;
QY 1 CTGACCCCTCTCGACTCGAGTTCG 28
DB 231 CTGGCCCATCCGACGCGAGAGTTCG 204

RESULT 7
AC021453/c 156165 bp DNA linear HTG 01-APR-2000
LOCUS Homo sapiens clone RP11-125C16, WORKING DRAFT SEQUENCE, 14
DEFINITION AC021453
AC021453.3 GI:7382318
VERSION

KEYWORDS HTG; HTGS PHASE1; HTGS_DRAFT.

SOURCE Homo sapiens (human)

ORGANISM Eukaryota Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 156165)

AUTHORS Biren,B., Linton,L., Nusbaum,C. and Lander,E.

TITLE Homo sapiens, clone RP11-125C16

JOURNAL Unpublished

REFERENCE 2 (bases 1 to 156165)

AUTHORS Biren,B., Linton,L., Nusbaum,C., Lander,E., Abraham,H., Allen,N., Anderson,S., Baldwin,J., Barna,N., Beckert,R., Beda,F., Boguslavsky,I., Bouhgalter,B., Brown,A., Burkett,G., Caetle,A., Chepel,Y., Colangelo,M., Collins,S., Collymore,A., Cooke,P., Dekrelano,K., Dewar,K., Domino,M., Doyle,M., Fensator,J., Ferreira,P., Fitzhugh,N., Forrest,C., Gage,D., Galagan,J., Gardyna,S., Grant,G., Hagos,B., Heatord,A., Horton,L., Howland,J.C., Johnson,R., Jones,C., Kann,L., Karatas,A., Klein,J., Landers,T., Lehoczy,J., Levine,R., Liu,G., Locke,K., Macdonald,P., Marguis,N., McEwan,P., McGuirk,A., McKernan,K., McHeeters,R., Meldrum,J., Meneue,L., Morrow,J., Naylor,J., Notman,C.H., O'Connor,T., O'Donnell,P., Ollivar,T.M., Peterson,K., Piere,N., Pisani,C., Pollara,V., Raymond,C., Riley,R., Rothman,D., Roy,A., Santos,R., Severi,P., Spencer,B., Stange-Thomann,N., Stojanovic,N., Subramanian,A., Talamas,J., Testaye,S., Theodore,J., Titrrell,A., Vassiliev,H., Viel,R., Vo,A., Wu,X., Wyman,D., Ye,W.J., Zimmer,A. and Zody,M.

Direct Submission

Submitted (16-JAN-2000) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA

On Apr 1, 2000 this sequence version replaced gi:6721267.

All repeats were identified using RepeatMasker:

Smit, A.F.A. & Green, P. (1996-1997)

http://ftp.genome.washington.edu/RM/RepeatMasker.html

Genome Center

Center: Whitehead Institute/ MIT Center for Genome Research

Center code: WIBR

Web site: http://www-seq.wi.mit.edu

Contact: sequence_submissions@genome.wi.mit.edu

Project Information

Center project name: L5169

Center clone name: 125.C.16

Summary Statistics

Sequencing vector: M13; M77815; 100% of reads

Chemistry: Dye-terminator Big Dye; 100% of reads

Assembly program: Phrap; version 0.960731

Consensus quality: 145142 bases at least Q40

Consensus quality: 151009 bases at least Q30

Consensus quality: 153345 bases at least Q20

Insert size: 160000; agarose-fp

Insert size: 154865; sum-of-coverage

Quality coverage: 4.1 in Q20 bases; agarose-fp

Quality coverage: 4.2 in Q20 bases; sum-of-coverage

NOTE: This is a 'working draft' sequence. It currently consists of 14 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown. This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

1 1836: contig of 1836 bp in length

1837 1936: gap of 100 bp

1937 5796: contig of 3860 bp in length

5797 5896: gap of 100 bp

5897 9797: contig of 3901 bp in length

9798 9897: gap of 100 bp

9898 13913: contig of 4022 bp in length

13920 14019: gap of 100 bp

14020 17400: contig of 3381 bp in length

17401 17500: gap of 100 bp

17501 21253: contig of 3753 bp in length

FEATURES

source

1. 156165

/organism="Homo sapiens"

/mol_type="genomic DNA"

/db_xref="taxon:9606"

/clone="RP11-125C16"

/clone_id="RP11-125C16"

1. 1836

/note="assembly_fragment"

1937. 5796

/note="assembly_fragment"

5897. 9797

/note="assembly_fragment"

9898. 13919

/note="assembly_fragment"

14020. 17400

/note="assembly_fragment"

17501. 21253

/note="assembly_fragment"

clone end:SP6

vector end:right"

21354. 31096

/note="assembly_fragment"

31197. 41465

/note="assembly_fragment"

41566. 50901

/note="assembly_fragment"

51002. 62041

/note="assembly_fragment"

62142. 75247

/note="assembly_fragment"

75348. 89874

/note="assembly_fragment"

89974. gap of 100 bp

116362: contig of 26388 bp in length

116363 116463: gap of 100 bp

156165: contig of 37503 bp in length.

Location/Qualifiers

1. 156165

21353: gap of 100 bp

21354 31096: contig of 9743 bp in length

31097 41465: contig of 10269 bp in length

41466 41565: gap of 100 bp

41566 50901: contig of 9336 bp in length

51001: gap of 100 bp

51002 62041: contig of 11040 bp in length

62042 62141: gap of 100 bp

62142 75247: contig of 13106 bp in length

75248 75347: gap of 100 bp

75348 89874: contig of 14527 bp in length

89875 89974: gap of 100 bp

116362: contig of 26388 bp in length

116363 116463: gap of 100 bp

156165: contig of 37503 bp in length.

ORIGIN

Query Match 66.7%; Score 20; DB 2; Length 156165;

Best Local Similarity 82.1%; Pred. No. 2.9e+02;

Matches 23; Conservative 0; Mismatches 5; Indels 0;

2 TGGACCCCTCTCGACTCGAGAGTTCCGC 29

DB 11285 TGGACCTCTCTCTATTCAGAGTTCTGC 11258

RESULT 8

LOCUS AC100852 157980 bp DNA linear PRI 29-AUG-2002

DEFINITION Homo sapiens chromosome 17, clone RP11-125C16, complete sequence.

ACCESSION AC100852

VERSION AC100852.2 GI:22539166

KEYWORDS HTG.

SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
AUTHORS Birren,B., Nussbaum,C. and Lander,E.
TITLE 1 (bases 1 to 157980)
JOURNAL Homo sapiens chromosome 17, clone RP11-125C16
REFERENCE Unpublished
AUTHORS 2 (bases 1 to 157980)
Birren,B., Linton,L., Nussbaum,C., Lander,E., Ali,A., Allen,N.,
Anderson,S., Barna,N., Bastien,V., Boguslavsky,L., Bouckgatter,B.,
Brown,A., Camarata,J., Campolano,A., Chang,J., Chazaro,B.,
Chapel,Y., Colangelo,M., Collins,S., Collymore,A., Cook,A.,
Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S., Dodge,S.,
Farooq,S., Fitzhugh,W., Gage,D., Galagan,J., Gargana,S.,
Ginde,S., Gord,S., Goyette,M., Graham,L., Grand-pierre,N.,
Hagos,B., Heatford,A., Horton,L., Hulme,W., Iliev,I., Johnson,R.,
Jones,C., Kamat,A., Karatas,A., Kells,C., Lakoque,K.,
Lamazares,R., Landers,T., Lehocsky,J., Levine,R., Matthews,C.,
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Norbu,C., Norman,C.H., O'Connor,T., O'Donnell,P., O'Neill,D.,
Oliver,J., Peterson,K., Phunkhang,P., Pierre,N., Pollara,V.,
Raymond,C., Retta,R., Rieback,M., Riley,R., Rise,C., Rogov,P.,
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Seaman,S., Severy,P., Spencer,B., Stange-Thomann,N., Stojanovic,N.,
Strauss,N., Subramaniam,A., Talamas,J., Testave,S., Theodore,J.,
Topham,K., Travers,M., Travis,N., Trigglio,J., Vassiliev,H.,
Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J., Young,G.,
Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.
TITLE Direct Submission
JOURNAL Submitted (22-NOV-2001) Whitehead Institute/MIT Center for Genome
REFERENCE Research, 320 Charles Street, Cambridge, MA 02141, USA
AUTHORS 3 (bases 1 to 157980)
Birren,B., Nussbaum,C., Lander,E., Ali,A., Allen,N., Anderson,S.,
Barna,N., Bastien,V., Bloom,T., Boguslavsky,L., Bouckgatter,B.,
Camarata,J., Chang,J., Chazaro,B., Chappel,Y., Collymore,A.,
Cook,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S., Dodge,S.,
Farooq,S., Ferreira,P., Fitzgerald,M., Gage,D., Galagan,J.,
Gargana,S., Gord,S., Graham,L., Grand-pierre,N., Hagos,B.,
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Karatas,A., Kells,C., Landers,T., Levine,R., Lindblad-Toh,K.,
Liu,G., Maclean,C., Macdonald,P., Major,J., Mathews,C.,
McCarthy,M., Meidrim,J., Meneus,L., Mihova,T., Mlenga,V.,
Murphy,T., Naylor,J., Nguyen,C., Nicol,R., Norbu,C., Norman,C.H.,
O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J., Peterson,K.,
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Smith,C., Spencer,B., Stange-Thomann,N., Stojanovic,N., Talamas,J.,
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Zembek,L., Zimmer,A. and Zody,M.
TITLE Direct Submission
JOURNAL Submitted (29-AUG-2002) Whitehead Institute/MIT Center for Genome
COMMENT Research, 320 Charles Street, Cambridge, MA 02141, USA
On Aug 29, 2002 this sequence version replaced gj:11048222.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html
----- Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WITR
Web site: http://www-seg.wi.mit.edu
Contact: sequence_submissions@genome.wi.mit.edu
----- Project Information
Center Project name: L21587
Center Clone name: 125_C_16

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Best Local Similarity 82.1%; Pred. No. 2.9e+02;
Matches 23; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

Qy      2 TGGACCCCTCTCGACTCGAGAGTCCGC 29
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RESULT 9
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LOCUS      Rattus norvegicus clone CH230-92124, WORKING DRAFT SEQUENCE, 4
DEFINITION      Rattus norvegicus (Norway rat)
ACCESSION      AC123011
VERSION      AC123011.3 GI:23665280
KEYWORDS      HTG; HTGS PHASE1; HTGS DRAFT; HTGS FULLTOP.
SOURCE      Rattus norvegicus (Norway rat)
ORGANISM      Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
Rattus.
REFERENCE      1 (bases 1 to 257595)

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AUTHORS

Muzny, D. Marie, Metzker, M. Lee, Abramson, S., Adams, C., Alder, J., Allen, C., Allen, H., Albrooks, S., Amin, A., Angiano, D., Anyalebechi, V., Aoyagi, A., Ayodeji, M., Bacc, E., Baden, H., Baldwin, D., Bandaranaike, D., Barber, M., Barnstead, M., Bernhamed, F., Biewald, K., Blair, J., Blankenburg, K., Blyth, P., Brown, M., Bryant, N., Buhay, C., Burch, P., Burrell, K., Calderon, E., Cardenas, V., Carter, K., Cavazos, I., Caesar, H., Center, A., Chacko, J., Chavez, D., Chen, G., Chen, R., Chen, Y., Chen, Z., Chu, J., Cleveland, C., Cockrell, R., Cox, C., Coyle, M., Cree, A., D'Souza, L., Davila, M. L., Davis, C., Davy-Carroll, L., De Anda, C., Dederich, D., Delgado, O., Denon, S., Deramo, C., Ding, Y., Dinh, H., Divya, K., Draper, H., Dugan-Rocha, S., Dunn, A., Durbin, K., Duval, B., Evans, K., Egan, A., Escotto, M., Eugene, C., Evans, C. A., Falls, T., Fan, G., Fernandez, S., Finley, M., Flagg, N., Forbes, L., Foster, M., Foster, P., Fraser, C. M., Gabisi, A., Ganta, R., Garcia, A., Garner, T., Garza, M., Gebregeorgis, E., Geer, K., Gill, R., Grady, M., Guerra, W., Guevara, M., Gunaratne, P., Haaland, W., Hamill, C., Hamilton, C., Hamilton, K., Harvey, Y., Havlak, P., Hawes, A., Henderson, N., Hernandez, J., Hernandez, R., Hines, S., Hladun, S. L., Hodgson, A., Hognes, M., Hollins, B., Howells, S., Hulys, S., Hume, J., Idlebird, D., Jackson, A., Jackson, L., Jacob, L., Jiang, H., Johnson, B., Johnson, R., Jolivet, A., Karpachy, S., Kelly, S., Kelly, S., Khan, Z., King, L., Kovar, C., Kowis, C., Kraft, C. L., Lebow, H., Levan, J., Lewis, L., Li, Z., Liu, J., Liu, J., Liu, W., Liu, Y., London, P., Longacre, S., Lopez, J., Lorenz, H., Louisa, L., Louisa, H., Lozano, R. J., Lu, X., Ma, J., Maheshwari, M., Mahindaratne, M., Mahmood, M., Malloy, K., Mangum, A., Mangum, B., Nagua, P., Martin, K., Martin, R., Martinez, E., Mawhiney, S., McLeod, M. P., McNeill, T. Z., Meenen, E., Milosavljevic, A., Miner, G., Minja, G., Montemayor, J., Moore, S., Morgan, M., Morris, K., Morris, S., Munidasa, M., Murphy, M., Nair, L., Nankervis, C., Neal, D., Newton, N., Nguyen, N., Norris, S., Nwankwem, O., Okunnu, G., Olajunsegun, A., Pal, S., Parks, K., Pasternak, S., Paul, H., Perez, A., Perez, L., Plannkoch, C., Plapper, F., Polidexter, A., Popovic, D., Primus, E., Pu, L. L., Puzo, M., Quiroz, J., Rachlin, E., Reeves, K., Regier, M. A., Reigh, R., Reilly, B., Reilly, M., Ren, Y., Reuter, M., Richards, S., Riggs, F., Rives, C., Rodkey, T., Rojas, A., Rose, M., Rose, R., Ruiz, S. J., Sanders, W., Savary, G., Scherer, S., Scott, G., Shatman, S., Shen, H., Shetty, J., Shvartsbeyn, A., Sisson, I., Sitter, C. D., Snajs, D., Sneed, A., Sodergren, E., Song, X. Z., Sorelle, R., Sosa, J., Steimle, M., Strong, R., Sutton, A., Szatek, A., Tabors, Z., Taylor, C., Taylor, T., Thomas, N., Thomas, S., Tingey, A., Trejos, P., Usaml, K., Valas, R., Vera, V., Villasana, D., Waldron, L., Walker, B., Wang, J., Wang, Q., Wang, S., Warren, J., Warren, R., Wei, X., White, F., Williams, G., Willson, R., Wleczky, R., Wooden, H., Worley, K., Wright, D., Wright, R., Wu, J., Yakub, S., Yen, J., Yoon, L., Yoon, V., Yu, F., Zhang, J., Zhou, J., Zhou, X., Zhao, S., Dunn, D., von Niederhausern, A., Weiss, R., Smith, D. R., Holt, R. A., Smith, H. O., Weinstock, G. and Gibbs, R. A.

TITLE

JOURNAL
REFERENCE
AUTHORS
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JOURNAL

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AUTHORS
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AUTHORS
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JOURNAL

Submitted (12-OCT-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
On Oct 10, 2002 this sequence version replaced gi:21909149.
The sequence in this assembly is a combination of BAC based reads and whole genome shotgun sequencing reads assembled using Atlas (http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig described in the feature table below represents a scaffold in the Atlas assembly (a 'contig-scaffold'). Within each contig-scaffold, individual sequence contigs are ordered and oriented, and separated by sized gaps filled with Ns to the estimated size. The sequence may extend beyond the ends of the clone and there may be sequence contigs within a contig-scaffold that consist entirely of whole


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Qy 2 TGAGCCCTCGACTCGAGAGTTCGCT 30
 Db 120906 TGAGCTACTCTGCTCCAGACTTCTCT 120878

RESULT 14
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 SEQUENCE, 4 unordered pieces.
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 VERSION AP006440.1 GI:30962586
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 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 REFERENCE
 1 Hattori,M., Ishii,K., Toyoda,A., Taylor,T.D., Hong-Seog,P.,
 Fujiyama,A., Yada,T., Tokoki,Y., Watanabe,H. and Sakaki,Y.
 Homo sapiens genomic DNA of 11q
 Published Only in Database (2003)
 2 (bases 1 to 189269)
 Hattori,M., Ishii,K., Toyoda,A., Taylor,T.D., Hong-Seog,P.,
 Fujiyama,A., Yada,T., Tokoki,Y., Watanabe,H. and Sakaki,Y.
 Direct Submission
 Submitted (19-MAY-2003) Masahira Hattori, The Institute of Physical
 and Chemical Research (RIKEN), Genomic Sciences Center (GSC),
 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
 (E-mail:hattori@gs.c.riken.go.jp, URL:http://hgp.gs.c.riken.go.jp/,
 Tel:81-45-503-9111, Fax:81-45-503-9170)

once, or longer because we provide a small overlap between
neighboring submissions.

FEATURES
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CRNA

CRNA

CRNA

ORIGIN

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Matches 23; Conservative 0; Mismatches 6; Indels 0; Gaps 0;

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RESULT 12

AB016816_6/c
WPCOMMENT

Sequence split into 9 fragments LOCUS AB016816 Accession AB016816

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AB016816_2	200001	310000
AB016816_3	200001	310000
AB016816_4	400001	410000
AB016816_5	500001	610000
AB016816_6	600001	710000
AB016816_7	700001	810000
AB016816_8	800001	907057

Continuation (7 of 9) of AB016816 from base 600001 (AB016816 Eremothecium gossypii ATCC

Query Match 64.7% Score 19.4; DB 8; Length 110000;
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Matches 23; Conservative 0; Mismatches 6; Indels 0; Gaps 0;

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RESULT 13
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LOCUS AC104791 159969 bp DNA linear PRI 21-FEB-2002

DEFINITION Homo sapiens BAC clone RP11-181K12 from 4, complete sequence.
ACCESSION AC104791 AC032008
VERSION AC104791.3 GI:18482313

KEYWORDS
HTG.
Homo sapiens (human)

SOURCE

ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.

REFERENCE
1 (bases 1 to 159969)
Toward a complete human genome sequence

JOURNAL
Genome Res. 8 (11), 1097-1108 (1998)

MEDLINE
99063792

PUBMED
9847074

REFERENCE
2 (bases 1 to 159969)
Isak, A., Meyer, R. and Creason, K.

REFERENCE
AUTHORS

TITLE
JOURNAL
REFERENCE
AUTHORS
TITLE
JOURNAL
Submitted (21-DEC-2001) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
4 (bases 1 to 159969)
Waterston, R.H.
Direct Submission
Submitted (03-FEB-2002) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
5 (bases 1 to 159969)
Waterston, R.
Direct Submission
Submitted (21-FEB-2002) Department of Genetics, Washington
University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
On Feb 3, 2002 this sequence version replaced gi:18042374.

COMMENT

Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: <http://genome.wustl.edu/gsc>
Contact: sapiens@watson.wustl.edu
----- Summary Statistics
Center project name: H_NH0181K12
Drafting Center: W18R

NOTICE: This sequence may not represent the entire insert of this
clone. It may be shorter because we only sequence overlapping
clone sections once, or longer because we provide a small overlap
between neighboring data submissions.

This sequence was finished as follows unless otherwise noted:
all regions were double stranded, sequenced with an alternate
chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such
as compressions and repeats; all regions were covered by sequence
from more than one subclone; and the assembly was confirmed by
restriction digest.

MAPPING INFORMATION:
Mapping information for this clone was provided by Dr. John D.
McPherson, Department of Genetics, Washington University, St. Louis
MO. For additional information about the map position of this
sequence, see <http://genome.wustl.edu/gsc>

SOURCE INFORMATION:
The RP11-11 human BAC library was made from the blood of one male
donor, as described by Osogawa, K., Moon, P.Y., Zhao, B., Frengen, E.,
Tateno, M., Catanesse, J.J. and de Jong, P.J. (1998) An improved
approach for construction of bacterial artificial chromosome
libraries. Genomics 51:1-8. The clone may be obtained either from
Research Genetics, Inc. (<http://www.resgen.com>) or Pieter de Jong
and coworkers at <http://www.chori.org>
VECTOR: pBAC3.6

NEIGHBORING SEQUENCE INFORMATION:
The clone sequenced to the left is RP11-292M9, the clone sequenced
to the right is RP11-203B7. Actual start of this clone is at base
position 1 of RP11-181K12; actual end is at base position 159969 of
RP11-181K12.

A transposon has been identified in the vector of this clone.

The sequence of AC032008 has been incorporated into AC104791.

FEATURES
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9849: gap of 100 bp
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10739 10838: gap of 100 bp
10839 11725: contig of 887 bp in length
11726 11825: gap of 100 bp
11825 12707: contig of 882 bp in length
12707 12807: gap of 100 bp
12807 13697: contig of 890 bp in length
13697 13797: gap of 100 bp
13797 14668: contig of 871 bp in length
14668 14768: gap of 100 bp
14768 15648: contig of 880 bp in length
15648 15748: gap of 100 bp
15748 16622: contig of 874 bp in length
16622 16722: gap of 100 bp
16722 17587: contig of 865 bp in length
17587 17687: gap of 100 bp
17687 18558: contig of 871 bp in length
18558 18658: gap of 100 bp
18658 19534: contig of 876 bp in length
19534 19634: gap of 100 bp
19634 20510: contig of 876 bp in length
20510 20610: gap of 100 bp
20610 21517: contig of 907 bp in length
21517 21617: gap of 100 bp
21617 22499: contig of 882 bp in length
22499 22599: gap of 100 bp
22599 23480: contig of 881 bp in length
23480 23580: gap of 100 bp
23580 24444: contig of 864 bp in length
24444 24544: gap of 100 bp
24544 25479: contig of 935 bp in length
25479 25579: gap of 100 bp
25579 26492: contig of 913 bp in length
26492 26592: gap of 100 bp
26592 27454: contig of 862 bp in length
27454 27554: gap of 100 bp
27554 28431: contig of 877 bp in length
28431 28531: gap of 100 bp
28531 29495: contig of 964 bp in length
29495 29595: gap of 100 bp
29595 30489: contig of 894 bp in length
30489 30589: gap of 100 bp
30589 31435: contig of 846 bp in length
31435 31535: gap of 100 bp
31535 32425: contig of 890 bp in length
32425 32525: gap of 100 bp
32525 33401: contig of 876 bp in length
33401 33501: gap of 100 bp
33501 34376: contig of 875 bp in length
34376 34476: gap of 100 bp
34476 35355: contig of 879 bp in length
35355 35455: gap of 100 bp
35455 36370: contig of 915 bp in length
36370 36470: gap of 100 bp
36470 37358: contig of 888 bp in length
37358 37458: gap of 100 bp
37458 38332: contig of 874 bp in length
38332 38432: gap of 100 bp
38432 39285: contig of 853 bp in length
39285 39385: gap of 100 bp
39385 40317: contig of 932 bp in length
40317 40417: gap of 100 bp
40417 41314: contig of 897 bp in length
41314 41414: gap of 100 bp
41414 42297: contig of 883 bp in length
42297 42397: gap of 100 bp
42397 43270: contig of 873 bp in length
43270 43370: gap of 100 bp
43370 44268: contig of 898 bp in length
44268 44368: gap of 100 bp
44368 45233: contig of 865 bp in length

45234 45333: gap of 100 bp
45333 46202: contig of 869 bp in length
46202 46302: gap of 100 bp
46302 47193: contig of 891 bp in length
47193 47293: gap of 100 bp
47293 48182: contig of 889 bp in length
48182 48282: gap of 100 bp
48282 49146: contig of 864 bp in length
49146 49246: gap of 100 bp
49246 50116: contig of 870 bp in length
50116 50216: gap of 100 bp
50216 51112: contig of 896 bp in length
51112 51212: gap of 100 bp
51212 52102: contig of 890 bp in length
52102 52202: gap of 100 bp
52202 53064: contig of 862 bp in length
53064 53164: gap of 100 bp
53164 54024: contig of 860 bp in length
54024 54124: gap of 100 bp
54124 54957: contig of 833 bp in length
54957 55057: gap of 100 bp
55057 55938: contig of 881 bp in length
55938 56038: gap of 100 bp
56038 56932: contig of 894 bp in length
56932 57032: gap of 100 bp
57032 57916: contig of 884 bp in length
57916 58016: gap of 100 bp
58016 58875: contig of 859 bp in length
58875 58975: gap of 100 bp
58975 59844: contig of 869 bp in length
59844 59944: gap of 100 bp
59944 60795: contig of 851 bp in length
60795 60895: gap of 100 bp
60895 61774: contig of 879 bp in length
61774 61874: gap of 100 bp
61874 62739: contig of 865 bp in length
62739 62839: gap of 100 bp
62839 63807: contig of 968 bp in length
63807 63907: gap of 100 bp
63907 64783: contig of 876 bp in length
64783 64883: gap of 100 bp
64883 65767: contig of 884 bp in length
65767 65867: gap of 100 bp
65867 66765: contig of 898 bp in length
66765 66865: gap of 100 bp
66865 67730: contig of 865 bp in length
67730 67830: gap of 100 bp
67830 68721: contig of 891 bp in length
68721 68821: gap of 100 bp
68821 69711: contig of 890 bp in length
69711 69811: gap of 100 bp
69811 70699: contig of 888 bp in length
70699 70799: gap of 100 bp
70799 71679: contig of 880 bp in length

Query Match 64.7%; Score 19.4; DB 2; Length 186472;
Best Local Similarity 79.3%; Pred. No. 5.3e+02;
Matches 23; Conservative 0; Mismatches 6; Indels 0; Gaps 0;
QY 1 CTGACCCCTCTCGACTCGAGAGTCCGC 29
Db 6946 CTGACCCCTCGTGTCTGAGAGTCCGC 6918

Search completed: April 26, 2005, 11:39:38
Job time : 1686 secs

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: April 26, 2005, 06:19:21 ; Search time 427 Seconds
(without alignments)
415.907 Million cell updates/sec

Title: US-10-086-062-4
Perfect score: 30
Sequence: 1 ctgaccctctcgcacgcagagctccgct 30

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 4390206 seqs, 2359870667 residues
Total number of hits satisfying chosen parameters: 8780412

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : N_Geneseq_16Dec04:*

- 1: geneseq1980s:*
- 2: geneseq1990s:*
- 3: geneseq2000s:*
- 4: geneseq2001as:*
- 5: geneseq2001bs:*
- 6: geneseq2002as:*
- 7: geneseq2002bs:*
- 8: geneseq2003as:*
- 9: geneseq2003bs:*
- 10: geneseq2003cs:*
- 11: geneseq2003ds:*
- 12: geneseq2004as:*
- 13: geneseq2004bs:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	30	100.0	30	6	AAD24069
2	19.2	64.0	1500	5	AAS11024
3	19	63.3	121124	12	ADQ97107
4	18.8	62.7	2481	10	ABT41766
5	18.8	62.7	23107	9	ADA02762
6	18.8	62.7	23107	10	ADB72500
7	18.8	62.7	23107	10	ADC85242
8	18.8	62.7	23107	12	ADM74357
9	18.6	62.0	3210	2	AAZ27624
10	18.6	62.0	3459	2	AAZ27623
11	18.4	61.3	319608	3	AAS151601
12	18.4	61.3	319608	5	AAS09301
13	18.2	60.7	340449	8	AAI52198
14	18	60.0	42325	10	ADM74382
15	18	60.0	110000	10	ADM74382
16	17.8	59.3	165	10	ADD49400
17	17.8	59.3	254	11	ADG00322
18	17.8	59.3	369	12	ABD11207
19	17.8	59.3	426	10	ADD49385
20	17.8	59.3	449	10	ADD49343

21	17.8	59.3	449	10	ADD49294	ADD49294 Human lun
22	17.8	59.3	986	6	ABQ46720	Abq46720 Oligonuc1
23	17.8	59.3	986	6	ABQ46721	Abq46721 Oligonuc1
24	17.8	59.3	1553	12	ADL12873	ADL12873 Human ste
25	17.8	59.3	1603	4	AAK77077	AAK77077 Human lmm
26	17.8	59.3	1731	11	ACN91401	ACN91401 Breast ca
27	17.8	59.3	2780	13	ADR25834	ADR25834 Breast ca
28	17.8	59.3	152141	8	ACA64961	ACA64961 Human BCR
29	17.8	59.3	153995	13	ABD33534	ABD33534 Murine ca
30	17.6	58.7	246	12	ADQ52469	ADQ52469 Human met
31	17.6	58.7	8119	3	AAZ35392	AAZ35392 Maize etca
32	17.6	58.7	49243	4	ABL03188	ABL03188 Drosophila
33	17.4	58.0	65	6	ABN51822	ABN51822 Mouse sp1
34	17.4	58.0	300	2	AAZ13036	AAZ13036 Human gen
35	17.4	58.0	300	2	AAZ98464	AAZ98464 Human can
36	17.4	58.0	411	4	AAI85652	AAI85652 Human pol
37	17.4	58.0	460	9	ACH41661	ACH41661 Human foe
38	17.4	58.0	594	12	ACH74968	ACH74968 Human gen
39	17.4	58.0	699	2	AAZ15926	AAZ15926 Human gen
40	17.4	58.0	729	12	ADQ63441	ADQ63441 Transcrip
41	17.4	58.0	778	6	ABK30430	ABK30430 Human G-p
42	17.4	58.0	936	13	ADR32877	ADR32877 Novel S.
43	17.4	58.0	949	13	ADS50577	ADS50577 Bacterial
44	17.4	58.0	1210	4	AAI41038	AAI41038 CDNA enco
45	17.4	58.0	1210	4	AAI34819	AAI34819 CDNA enco

ALIGNMENTS

RESULT 1	
ID	AAD24069 standard; DNA; 30 BP.
XX	
AC	AAD24069;
XX	
DT	09-APR-2002 (first entry)
XX	
DE	Maize engineered Ubi-1 promoter heat shock element #3.
XX	
KW	Gene expression; maize; ubiquitin promoter; Ubi-1; HSE;
KM	heat shock element; agronomic gene; ds.
XX	
OS	Zea mays.
XX	
FT	Key
FT	Location/Qualifiers
FT	1.15
FT	/*tag= a
FT	/note= "5' heat shock element"
FT	16..30
FT	/*tag= b
FT	/note= "3' heat shock element"
XX	
PN	W0200194394-A2.
XX	
PD	13-DEC-2001.
XX	
PE	08-JUN-2001; 2001WO-US018689.
XX	
PR	09-JUN-2000; 2000US-00590558.
XX	
PA	(PROD-) PRODIGENE INC.
XX	
PI	Jilka JM, Hood EE, Howard JA;
XX	
DR	WPI; 2002-122117/16.
XX	
PT	New promoter sequences for causing expression of a structural gene
PT	especially agronomic gene or open reading frame in a plant cell,
XX	comprises engineered versions of the maize ubiquitin promoter.
PS	Claim 6; Page 54; 68pp; English.
XX	

CC The invention relates to a promoter sequence capable of directing
 CC expression of a nucleotide sequence in a plant cell, comprising maize
 CC ubiquitin (ubi-1) promoter sequence with a modification so that it does
 CC not include two overlapping heat shock elements (HSE) or it directs
 CC expression to increase the endosperm/embryo expression ratio of the
 CC protein when compared to the ratio from a wild-type ubiquitin promoter.
 CC The modified ubi-1 promoter comprises a deletion of 3', 5' or both HSEs,
 CC two non-overlapping/adjacent HSEs, replacement of HSEs with a trimer of a
 CC seed specific element from the promoter of pea lectin gene Psi, or
 CC insertion of a transcription factor binding site in the HSE region. An
 CC expression construct comprising modified ubi-1 promoter is useful for
 CC causing expression of a structural gene (agronomic genes) or open reading
 CC frame in a plant cell. The modified ubi-1 promoter increases expression
 CC levels beyond those observed with native ubiquitin promoter. The present
 CC sequence is maize engineered ubi-1 promoter with heat shock elements
 CC adjacent placed. Note: The present sequence is also shown in claim 26,
 CC page 56 of the specification. However, this sequence has an additional
 CC nucleotide at the 3' end

Sequence 30 BP; 4 A; 12 C; 7 G; 7 T; 0 U; 0 Other;

Query Match 100.0%; Score 30; DB 6; Length 30;
 Best Local Similarity 100.0%; Pred. No. 0.0012;
 Matches 30; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 CTGACCCCTCTGACTCGAGATTCCGCT 30
 1 CTGACCCCTCTGACTCGAGATTCCGCT 30

RESULT 2
 AAS1024/c
 ID AAS1024 standard; DNA; 1500 BP.

AC AAS1024;
 DT 11-SEP-2003 (revised)
 DT 24-OCT-2001 (first entry)

DE Vibrio cholera 16S ribosomal RNA gene.

KM Antisense; bacterial 16S ribosomal RNA; rRNA; bacterial infection; human;
 KM food grain supplement; livestock; poultry; therapeutic; ds.

OS Vibrio cholerae.

PN WO200142457-A2.

PD 14-JUN-2001.

PF 29-NOV-2000; 2000WO-US042391.

PR 29-NOV-1999; 99US-0168150P.

PA (AVIB-) AVI BIOPHARMA INC.

PI Iversen PL;

DR WPI; 2001-457295/49.

PT Antibacterial compound, useful for treating bacterial infections and as
 PT livestock and poultry food supplement, comprises antisense
 PT oligonucleotides complementary to bacterial 16S and 23S rRNA.

PS Disclosure; Page; 62pp; English.

CC AAS1021-AAS1024 represent the coding sequences of bacterial 16S
 CC ribosomal RNA (rRNA) genes. The sequences were used to design anti-
 CC bacterial compounds comprising substantially uncharged antisense
 CC oligomers containing 8-40 nucleotide subunits, including a targeting
 CC nucleic acid sequence at least 10 nucleotides in length which is
 CC complementary to a bacterial 16S or 23S rRNA nucleic acid sequence. The
 CC antisense oligomers are used for treating a bacterial infection in a

CC human or a mammalian animal produced by *Escherichia coli*, *Salmonella*
 CC typhimurium, *Pseudomonas aeruginosa*, *Vibrio cholerae*, *Neisseria*
 CC gonorrhoea, *Helicobacter pylori*, *Bartonella henselae*, *Haemophilus*
 CC influenzae, *Shigella dysenteriae*, *Staphylococcus aureus*, *Mycobacterium*
 CC tuberculosis, *Streptococcus pneumoniae*, *Treponema pallidum* and *Chlamydia*
 CC trachomatis. The antibacterial compound may be used as a food grain
 CC supplement in livestock and poultry food composition. Note: The present
 CC sequence is not shown in the specification but has been accessed from
 CC GenBank using the appropriate accession number given in the
 CC specification. (Updated on 11-SEP-2003 to standardise OS field)

Sequence 1500 BP; 376 A; 326 C; 482 G; 312 T; 0 U; 4 Other;

Query Match 64.0%; Score 19.2; DB 5; Length 1500;
 Best Local Similarity 75.0%; Pred. No. 1.1e+02;
 Matches 21; Conservative 2; Mismatches 5; Indels 0; Gaps 0;

OY 1 CTGACCCCTCTGACTCGAGATTCCG 28
 212 CTGGGCCCATCCCGACGCGARAGTCCG 185

RESULT 3
 ADQ97107/c
 ID ADQ97107 standard; DNA; 121124 BP.

AC ADQ97107;

DT 07-OCT-2004 (first entry)

DE Mouse cancer associated sequence MD08-002, SEQ ID 83.

KM Cytostatic; Gene Therapy; cancer; leukemia; lymphoma; Mouse; ds.

OS Mus musculus.

PN WO2004060304-A2.

PD 22-JUL-2004.

PF 22-DEC-2003; 2003WO-US041389.

PR 27-DEC-2002; 2002US-00330773.

PA (SAGR-) SAGRES DISCOVERY INC.

PI Morris DW, Malandro MS;

DR WPI; 2004-543781/52.

PT New isolated cancer associated nucleic acids comprising at least 10
 PT contiguous nucleotides, useful for diagnosing, preventing and/or treating
 PT cancers such as leukemia and lymphoma.

PS Claim 1; SEQ ID NO 83; 199pp; English.

CC The present invention relates to cancer associated sequences (ADQ97025-
 CC ADQ98004). The sequences are useful for the diagnosis, prevention and/or
 CC treatment of cancer, such as leukemia and lymphoma. Note: The sequence
 CC data for this patent did not form part of the printed specification, but
 CC was obtained in electronic format directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences.

Sequence 121124 BP; 32972 A; 25314 C; 26641 G; 35451 T; 0 U; 746 Other;

Query Match 63.3%; Score 19; DB 12; Length 121124;
 Best Local Similarity 81.5%; Pred. No. 1.9e+02;
 Matches 22; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

OY 4 GACCCCTCTGACTCGAGATTCCGCT 30
 DB 88764 GCCCCTCTGATTACAGATTCCGCT 88738

RESULT 4
ABT1766/C
ID ABT1766 standard; DNA; 2481 BP.
XX
AC ABT1766;
XX
DT 26-JUN-2003 (first entry)
XX
DE Toxicity modelling related rat gene SEQ ID NO 1468.
XX
KW Toxic effect; gene expression profile; renal toxicity; toxicity marker;
KW database; drug screening; toxicity assay; rat; ds.
XX
OS Rattus norvegicus.
XX
PN WO200295000-A2.
XX
PD 28-NOV-2002.
XX
PE 22-MAY-2002; 2002WO-US016173.
XX
PR 22-MAY-2001; 2001US-0292335P.
PR 13-JUN-2001; 2001US-0297523P.
PR 19-JUN-2001; 2001US-0298925P.
PR 10-JUL-2001; 2001US-0303807P.
PR 10-JUL-2001; 2001US-0303808P.
PR 10-JUL-2001; 2001US-0303810P.
PR 28-AUG-2001; 2001US-0315047P.
PR 27-SEP-2001; 2001US-0324928P.
PR 22-OCT-2001; 2001US-0330462P.
PR 01-NOV-2001; 2001US-0330867P.
PR 21-NOV-2001; 2001US-0331805P.
PR 06-DEC-2001; 2001US-0336144P.
PR 19-DEC-2001; 2001US-0340873P.
PR 21-FEB-2002; 2002US-0357842P.
PR 21-FEB-2002; 2002US-0357843P.
PR 21-FEB-2002; 2002US-0357844P.
PR 15-MAR-2002; 2002US-0364134P.
PR 08-APR-2002; 2002US-0370144P.
PR 08-APR-2002; 2002US-0370206P.
PR 17-APR-2002; 2002US-0372794P.
PR 21-APR-2002; 2002US-0371679P.
XX
PA (GENE-) GENE LOGIC INC.
XX
PI Mendrick D, Porter M, Johnson K, Higgs B, Castle A, Elashoff M;
XX
DR WPI; 2003-148464/14.
XX
FT Predicting at least one toxic effect of a compound, useful for toxicity
PT modelling, comprises preparing a gene expression profile of a tissue or
PT cell sample exposed to the compound, and comparing the gene expression
PT profile to a database.
XX
PS Example 4; Page; 446pp; English.
XX
XX The invention relates to a novel method of predicting at least one toxic
XX effect of a compound. The method comprises a gene expression profile of a
XX tissue or cell sample exposed to the compound, and comparing the gene
XX expression profile to a database comprising at least part of the data or
XX information given in the specification. The methods are useful for
XX predicting at least one toxic effect of a compound, predicting the
XX progression of a toxic effect of a compound, predicting the renal
XX toxicity of a compound, or identifying toxicity markers in tissues or
XX cells exposed to known renal toxin. The genes are useful as toxicity
XX markers in drug screening and toxicity assays, in monitoring disease or
XX physiological states, or disease progression. This polynucleotide
XX represents a rat DNA sequence relating to the toxic effect database
XX described in the specification. NOTE: The sequence data for this patent
XX did not form part of the printed specification, but was obtained in
XX electronic format directly from the World Intellectual Property

CC Organization
XX
SQ Sequence 2481 BP; 599 A; 653 C; 555 G; 674 T; 0 U; 0 Other;
CC
Query Match 62.7%; Score 18.8; DB 10; Length 2481;
Best Local Similarity 76.7%; Pred. No. 1.7e+02;
Matches 23; Conservative 0; Mismatches 7; Indels 0; Gaps 0;
QY 1 CTGACCCCTCTCGACTCGAGATTCCGCT 30
DB 1077 CTGTCCCTCTCTCGATTAGGAGTTCAAGT 1048
RESULT 5
ADA02762
ID ADA02762 standard; DNA; 23107 BP.
XX
AC ADA02762;
XX
DT 06-NOV-2003 (first entry)
XX
DE Human RUNX3 carcinoma associated gene, SEQ ID NO:1280.
XX
KW Human; carcinoma associated; oncogene; carcinoma; cancer; breast;
KW prostate; lymphoma; leukaemia; cytostatic; gene therapy; drug screening;
KW gene; ds.
XX
OS Homo sapiens.
XX
PN WO2003057146-A2.
XX
PD 17-JUL-2003.
XX
PE 26-DEC-2002; 2002WO-US041414.
XX
PR 26-DEC-2001; 2001US-00035832.
XX
PA (SAGR-) SAGRES DISCOVERY.
XX
PI Morris DW;
XX
DR WPI; 2003-587066/55.
XX
FT New recombinant nucleic acid encoding carcinoma associated protein,
PT useful for preparing compositions for treating carcinomas.
XX
PS Claim 1; SEQ ID NO 1280; 245pp; English.
XX
XX The invention relates to recombinant carcinoma associated (CA) nucleic
XX acid sequences from mouse and human (ADA01482-ADA03094), and to
XX recombinant carcinoma associated proteins (CAP) encoded by them. The
XX invention also encompasses expression vectors and host cells comprising a
XX CA nucleic acid, a polypeptide (especially an antibody) that specifically
XX binds to the protein, and a biochip comprising CA nucleic acid or
XX fragments thereof. The sequences of the invention were identified using
XX oncogenic retroviruses, which insert into the genome of the host organism
XX at random. Many of these do not carry transduced host oncogenes or
XX pathogenic trans-acting viral genes, meaning that cancer incidence is a
XX direct consequence of the effects of proviral integration into host
XX protooncogenes. The CA nucleic acid sequences can be used to diagnose
XX carcinoma (especially breast cancer, prostate cancer, lymphoma or
XX leukaemia) or a propensity to carcinoma by determination of the sequence
XX of a CA gene, or by determination of CA gene expression in particular
XX tissues. CA nucleic acids, proteins and antibodies are also useful as
XX therapeutic agents and in screening and evaluating drug candidates. The
XX present sequence represents a specifically claimed human CA nucleic acid
XX sequence of the invention. Note: The complete sequence data for this
XX patent did not form part of the printed specification, but was obtained
XX in electronic format directly from WIPO at
XX ftp.wipo.int/pub/published_pct_sequences.
SQ Sequence 23107 BP; 5308 A; 6264 C; 6229 G; 5306 T; 0 U; 0 Other;

Query Match 62.7%; Score 18.8; DB 9; Length 23107;
 Best Local Similarity 76.7%; Pred. No. 2.1e+02;
 Matches 23; Conservative 0; Mismatches 7; Indels 0; Gaps 0;

OY 1 CTGGACCCCTCTGACTCGAGATTCCGCT 30
 DB 14873 CTGGAGACCCCTGACTCCAGATTCCGCT 14902

RESULT 6

ADB72500
 ID ADB72500 standard; DNA; 23107 BP.

AC ADB72500;

DT 04-DEC-2003 (first entry)

DE Human Runx3 gene.

human; ds; cytosstatic; gene therapy; vaccine; carcinoma; lymphomas;
 cancer; neoplasm; adenocarcinoma; sarcoma; gene.

OS Homo sapiens.

PN MO2003008583-A2.

PD 30-JAN-2003.

PF 26-DEC-2001; 2001WO-US051291.

PR 02-MAR-2001; 2001US-00798586.

PR 23-OCT-2001; 2001US-00004113.

PR 08-NOV-2001; 2001US-00052482.

PR 30-NOV-2001; 2001US-00997722.

PR 20-DEC-2001; 2001US-00034650.

PA (SAGR-) SAGRES DISCOVERY.

PI Morris DW, Engelhard EK;

PT WPI; 2003-239337/23.

PS Claim 1; SEQ ID NO 328; 2304pp; English.

XX The invention relates to a novel recombinant nucleic acid comprising a

XX CC nucleotide sequence selected from any of the 660 sequences fully defined

XX CC in the specification. A polynucleotide of the invention has cytosstatic

XX CC activity, and may have a use in gene therapy, or in a vaccine. The

XX CC recombinant nucleic acids and polypeptides are useful for treating

XX CC carcinomas, e.g. lymphomas, cancers, neoplasm, adenocarcinoma, and

XX CC sarcomas. The present sequence represents a human gene of the invention.

XX Sequence 23107 BP; 5308 A; 6264 C; 6229 G; 5306 T; 0 U; 0 Other;

OY 1 CTGGACCCCTCTGACTCGAGATTCCGCT 30
 DB 14873 CTGGAGACCCCTGACTCCAGATTCCGCT 14902

RESULT 7

ADB72500
 ID ADB72500 standard; DNA; 23107 BP.

AC ADB72500;

DT 01-JAN-2004 (first entry)

XX Human Runx3 genomic sequence.

DE Cytostatic; gene therapy; vaccine; cancer; carcinoma-associated gene; CA;

KW secreted; transmembrane; intracellular; ds.

OS Homo sapiens.

PN MO2003045230-A2.

PD 05-JUN-2003.

PF 02-DEC-2002; 2002WO-US038582.

PR 30-NOV-2001; 2001US-00997722.

PA (SAGR-) SAGRES DISCOVERY.

PI Morris DW, Engelhard EK;

PT WPI; 2003-513603/48.

PS New recombinant nucleic acid comprising a nucleotide sequence of any of

XX the carcinoma-associated (CA) genes, useful for screening for drug

XX candidates for diagnosing or treating carcinomas.

XX Claim 1; SEQ ID NO 28; 983pp; English.

XX The invention relates to a recombinant nucleic acid comprising a

XX CC nucleotide sequence selected from any of the fully defined carcinoma-

XX CC associated (CA) genes from the 50 tables given in the specification. The

XX CC CA proteins are secreted, transmembrane or intracellular proteins. The

XX CC recombinant nucleic acids are useful for screening for drug candidates

XX CC for diagnosing or treating carcinomas. Sequences given in ADB72515-

XX ADB72514 represent CA genes of the invention.

XX Sequence 23107 BP; 5308 A; 6264 C; 6229 G; 5306 T; 0 U; 0 Other;

OY 1 CTGGACCCCTCTGACTCGAGATTCCGCT 30
 DB 14873 CTGGAGACCCCTGACTCCAGATTCCGCT 14902

ADMT74357
 ID ADMT74357 standard; DNA; 23107 BP.

XX ADMT74357;

DT 01-JUN-2004 (first entry)

DE Human carcinoma associated (CA) nucleic acid #13.

KW Human; carcinoma associated nucleic acid; CA nucleic acid; gene; ds;

KW carcinoma associated protein; CAP; carcinoma; leukemia; lymphoma;

XX cytosstatic.

OS Homo sapiens.

PN US2004072154-A1.

PD 15-APR-2004.

PF 30-NOV-2001; 2001US-00997722.

PR 22-DEC-2000; 2000US-00747377.

PR 02-MAR-2001; 2001US-00798586.

PA (MORR/) MORRIS D W.

PA	(ENSEB// ENGELHARD E K.
XX	
PI	Morris DW, Engelhard EK;
DR	WPI, 2004-328562/30.
XX	
PT	New carcinoma associated gene or protein, useful for preparing a
PR	composition for diagnosing or treating carcinoma e.g., leukemia or
PT	Lymphoma.
XX	
PS	Claim 1; SEQ ID NO 28; 29pp; English.
XX	
CC	The invention relates to new recombinant nucleic acids. The invention
CC	also relates to a host cell comprising a recombinant nucleic acid or
CC	expression vector, an expression vector comprising a recombinant nucleic
CC	acid, a recombinant protein, a method of screening for drug candidates, a
CC	method of screening for a bioactive agent capable of binding to a
CC	carcinoma associated protein (CAP) encoded by a nucleotide sequence, a
CC	method of screening for a bioactive agent capable of modulating the
CC	activity of a CAP, a method of evaluating the effect of a candidate
CC	carcinoma drug, a method of diagnosing carcinoma, a method for inhibiting
CC	the activity of a CAP, a method of treating carcinomas, a method of
CC	neutralising the effect of a CAP and a method of diagnosing carcinoma or
CC	propensity to carcinoma. A method of evaluating the effect of a candidate
CC	carcinoma drug comprises administering the drug to a patient, removing a
CC	cell sample from the patient and determining alterations in the
CC	expression or activation of a gene comprising the nucleotide sequence. A
CC	method of diagnosing carcinoma comprises determining the expression of
CC	one or more genes comprising the nucleic acid sequence in a first tissue
CC	type of a first individual and comparing the expression of the gene from
CC	a second normal tissue type from the first individual or a second
CC	unaffected individual, where a difference in the expression indicates
CC	that the first individual has carcinoma. A method of inhibiting the
CC	activity of a CAP comprises binding an inhibitor to the CAP. Treating
CC	carcinomas comprises administering to a patient an inhibitor of CAP.
CC	Neutralising the effect of a CAP comprises contacting an agent specific
EC	for the CAP. The polypeptide specifically binds to the protein encoded by
EC	the nucleic acid. It comprises an antibody that specifically binds to the
TC	protein encoded by the nucleic acid. The nucleic acids are useful for
CC	preparing a composition for diagnosing or treating carcinoma e.g.,
CC	leukemia or lymphoma. This sequence represents a human carcinoma
CC	associated (CA) nucleic acid of the invention. Note: The sequence data
CC	for this patent did not form part of the printed specification but was
CC	obtained in electronic format directly from USPRO at
CC	seqdata.uspto.gov/sequence.html.
XX	
SQ	Sequence 23107 BP; 5308 A; 6264 C; 6229 G; 5306 T; 0 U; 0 Other;
	Query Match 62.7%; Score 18.8; DB 12; Length 23107;
	Best Local Similarity 76.7%; Pred. No. 2.1e+02;
	Matches 23; Conservative 0; Mismatches 7; Indels 0; Gaps 0
OY	1 CTGACCCCCTCTCGACTCGAGAGTTCGGCT 30
Dd	
	14873 CTGGAGGCCCTCGACTCCAGAATTCGCT 14902
RESULT 9	
ID	AA227624/C
XX	AA227624 standard; DNA; 3210 BP.
XX	
XX	AA227624;
DT	20-DEC-1999 (first entry)
DE	
XX	Plasmid SPF-1.
XX	
KW	Extracellular compartment modification; floral cell; self-compatibility;
KV	pollen-pistil interaction; self-incompatibility; insect growth control;
KX	plasmid SPF-1, GRIS363 gene; cysteine protease inhibitor gene; ss.
XX	
OS	Synthetic.
XX	

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PN      WO9949063-A1.
XX      30-SEP-1999.
PD      19-MAR-1999;      99WO-CA000237.
XX      20-MAR-1998;      98US-0078728P.
PR      (MTAC ) CANADA MIN AGRIC & AGRI-FOOD CANADA.
XX      Robert LS,      Gleddie S;
XX      WPI, 1999-591104/50.
XX      protein expression in floral cells for peptide display, mediating plant
XX      sterility, and modifying pollen-pistil interactions.
XX      Example 6; Page 103-104; 113pp; English.
PS      This sequence represents the plasmid SPF-1, containing a fusion of the
CC      Brassica napus GPIS363 gene to the Onchocerca protease inhibitor gene.
CC      The invention relates to a method for modifying the extracellular
CC      compartment of a floral cell of a plant, that comprises expressing a
CC      construct comprising a gene of interest encoding a protein, fusion
CC      protein or peptide, or a fragment of them, which is capable of modifying
CC      the composition of the extracellular compartment of the floral cell and
CC      altering either the function, use or development of the floral cell or
CC      modifying the interaction of the floral cell with other cells, within an
CC      anther or pistil cell. The method is used to modify pollen-pistil
CC      interaction or function, which mediates, produces or prevents self-
CC      compatibility, self-incompatibility, out- or in-crossing or combinations
CC      of these. The method is also used for localizing proteins on the surface
CC      of pollen for the purpose of peptide display. The protein localized on
CC      the surface of the pollen may be an antibody or antigen or is a protein
CC      that is effective in controlling insect growth, behaviour, feeding,
CC      development or reproduction
SQ      Sequence 3210 BP, 1050 A; 626 C; 595 G; 939 T; 0 U; 0 Other;
Query Match      62.0%; Score 18.6; DB 2; Length 3210;
Best Local Similarity      84.0%; Pred. No. 2,2e+02;
Matches      21; Conservative      0; Mismatches      4; Indels      0; Gaps      0;
OY      1 CTGACCCCTCTCGACTCGAGAGTT 25
DB      1232 CTGACCCCTTTCGATCGATAGTT 1208
      |||||
RESULT 10
AAZ27623/C
ID      AAZ27623 standard; DNA; 3459 BP.
XX      AAZ27623;
AC      20-DEC-1999 (first entry)
DT      20-DEC-1999 (first entry)
XX      Plasmid SPF-1.
DE      Plasmid SPF-1.
XX      Extracellular compartment modification; floral cell; self-compatibility;
KM      pollen-pistil interaction; self-incompatibility; insect growth control;
KW      plasmid SPF-1; GPIS363 gene; cysteine protease gene; ss..
XX      Synthetic.
OS      Synthetic.
XX      WO9949063-A1.
XX      30-SEP-1999.
PD      19-MAR-1999;      99WO-CA000237.
XX      20-MAR-1998;      98US-0078728P.
PR      (MTAC ) CANADA MIN AGRIC & AGRI-FOOD CANADA.
XX      Robert LS,      Gleddie S;
XX      WPI, 1999-591104/50.
XX      protein expression in floral cells for peptide display, mediating plant
XX      sterility, and modifying pollen-pistil interactions.
XX      Example 6; Page 103-104; 113pp; English.
PS      This sequence represents the plasmid SPF-1, containing a fusion of the
CC      Brassica napus GPIS363 gene to the Onchocerca protease inhibitor gene.
CC      The invention relates to a method for modifying the extracellular
CC      compartment of a floral cell of a plant, that comprises expressing a
CC      construct comprising a gene of interest encoding a protein, fusion
CC      protein or peptide, or a fragment of them, which is capable of modifying
CC      the composition of the extracellular compartment of the floral cell and
CC      altering either the function, use or development of the floral cell or
CC      modifying the interaction of the floral cell with other cells, within an
CC      anther or pistil cell. The method is used to modify pollen-pistil
CC      interaction or function, which mediates, produces or prevents self-
CC      compatibility, self-incompatibility, out- or in-crossing or combinations
CC      of these. The method is also used for localizing proteins on the surface
CC      of pollen for the purpose of peptide display. The protein localized on
CC      the surface of the pollen may be an antibody or antigen or is a protein
CC      that is effective in controlling insect growth, behaviour, feeding,
CC      development or reproduction
SQ      Sequence 3210 BP, 1050 A; 626 C; 595 G; 939 T; 0 U; 0 Other;
Query Match      62.0%; Score 18.6; DB 2; Length 3210;
Best Local Similarity      84.0%; Pred. No. 2,2e+02;
Matches      21; Conservative      0; Mismatches      4; Indels      0; Gaps      0;
OY      1 CTGACCCCTCTCGACTCGAGAGTT 25
DB      1232 CTGACCCCTTTCGATCGATAGTT 1208
      |||||

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XX Robert LS, Gledlie S;
PI WPI; 1999-591104/50.
XX Protein expression in floral cells for peptide display, mediating plant
XX sterility, and modifying pollen-pistil interactions.
XX Example 5; Page 100-102; 113pp; English.
PS
XX This sequence represents the plasmid SPF-1, containing a fusion of the
XX Brassica napus GPIS363 gene to the Strophilus cysteine protease gene. The
XX invention relates to a method for modifying the extracellular compartment
XX of a floral cell of a plant, that comprises expressing a construct
XX comprising a gene of interest encoding a protein, fusion protein or
XX peptide, or a fragment of them, which is capable of modifying the
XX composition of the extracellular compartment of the floral cell and
XX altering either the function, use or development of the floral cell or
XX modifying the interaction of the floral cell with other cells, within an
XX anther or pistil cell. The method is used to modify pollen-pistil
XX interaction or function, which mediates, produces or prevents self-
XX compatibility, self-incompatibility, out- or in-crossing or combinations
XX of these. The method is also used for localizing proteins on the surface
XX of pollen for the purpose of peptide display. The protein localized on
XX the surface of the pollen may be an antibody or antigen or is a protein
XX that is effective in controlling insect growth, behaviour, feeding,
XX development or reproduction
XX
SQ Sequence 3459 BP; 1082 A; 721 C; 678 G; 978 T; 0 U; 0 Other;
- Query Match 62.0%; Score 18.6; DB 2; Length 3459;
- Best Local Similarity 84.0%; Pred. No. 2.2e+02;
- Matches 21; Conservative 0; Mismatches 4; Indels 0; Gaps 0;
OY 1 CTGAGCCCTCTCGACTCGAGACTT 25
Db 1232 CTGAGCCCTCTCGAGACTT 1208
-
RESULT 11
AAH51601
XX AAH51601 standard; DNA; 319608 BP.
AC
XX AAH51601;
XX
DT 29-AUG-2001 (first entry)
XX
DE Human chromosome 13q31-q33 genomic nucleotide sequence.
XX
KM sbg1; g3465; sbg2; g35017; g35018; chromosome 13q31-q33; haplotype;
XX biallelic marker; polymorphism; schizophrenia; bipolar disorder; ds.
XX
OS Homo sapiens.
XX
PN WO200058510-A2.
XX
PD 05-OCT-2000.
XX
PE 30-MAR-2000; 2000MO-IB000435.
XX
XX 30-MAR-1999; 99US-0126903P.
XX 30-APR-1999; 99US-0131971P.
XX 30-APR-1999; 99US-0132065P.
XX 14-JUL-1999; 99US-0143928P.
XX 27-JUL-1999; 99US-0145915P.
XX 29-JUL-1999; 99US-0146452P.
XX 29-JUL-1999; 99US-0146453P.
XX 28-OCT-1999; 99US-0162288P.
XX
PA (GEST ) GENSET.
XX
XX Cohen D, Blumenfeld W, Chumakov I, Bougueleret L, Bihain B,
PI Bastoux L;

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XX WPI; 2000-619082/59.
DR Polynucleotides comprising sequences from sbg1 and g35018 biallelic
XX markers are used for genotyping and detecting schizophrenia or bipolar
PT disorder and predisposition to these disorders.
XX
XX Claim 1; Page 409-493; 737pp; English.
PS
XX AAH51601 represents a human genomic nucleotide sequence comprising sbg1,
XX g3465, sbg2, g35017 and g35018 nucleic acid sequences located on the
XX human chromosome 13q31-q33 locus. The nucleotide sequences contain
XX biallelic markers and polymorphisms. Sequences AAH51602 - AAH51626 and
XX AAH62907 - AAH62915 represent cDNA human sbg1 cDNA sequences and protein
XX products. AAH51627 - AAH51631 and AAH62916 - AAH62918 represent g35018
XX cDNA sequences and protein products. Primers AAH51632 - AAH51699 are used
XX to isolate sbg1 cDNAs, while sbg1 exons from different primates are
XX represented by sequences AAH51642 - AAH51699. Nucleotide sequences of
XX amplicons which comprise biallelic markers located on the chromosome
XX 13q31-q33 locus are represented in AAH51700 - AAH51817. Biallelic markers
XX are represented in the sequences by degenerate/undefined base codes. PCR
XX primers AAH51818 and AAH51819 are used in the isolation of sequences of
XX the invention. The biallelic marker containing nucleotide sequences are
XX used to determine the identity of the nucleotide at a biallelic marker in
XX a sample DNA sequence. The nucleotide sequences may be labelled and used
XX for genotyping by determining the identity of a nucleotide at a Region D-
XX related biallelic marker in a biological sample from single or multiple
XX subjects. By determining the frequency of a biallelic marker in a
XX population an association between a genotype and a trait, a haplotype and
XX a trait and a phenotype and a trait can be detected. The sequences can be
XX used to determine a predisposition to or early onset of schizophrenia or
XX bipolar disorder or a beneficial response to or side effects related to
XX treatment against schizophrenia or bipolar disorder
XX
SQ Sequence 319608 BP; 101600 A; 56677 C; 58335 G; 102722 T; 0 U; 274 Other;
- Query Match 61.3%; Score 18.4; DB 3; Length 319608;
- Best Local Similarity 78.6%; Pred. No. 3.8e+02;
- Matches 22; Conservative 0; Mismatches 6; Indels 0; Gaps 0;
OY 1 CTGAGCCCTCTCGACTCGAGACTTCCG 28
Db 8690 CTGAGCCCTCTCGACTTGAATTGACG 8717
-
RESULT 12
AAS09301
XX AAS09301 standard; DNA; 319608 BP.
AC
XX AAS09301;
XX
DT 26-SEP-2001 (first entry)
XX
DE Human schizophrenia associated gene g35030 and biallelic markers A1-A71.
XX
XX Human: g35030; biallelic marker; A1-A71; chromosome 13q31-q33;
XX schizophrenia; bipolar disorder; ds.
XX
OS Homo sapiens.
XX
XX
FH Key Location/Qualifiers
FH primer_bind 7938..7958
FT /tag= a
FT /note= "binds primer 99-27943.rp"
FT primer_bind 8297..8315
FT /tag= b
FT /note= "binds primer 99-27943-150.mis"
FT misc_binding 8304..8328
FT /tag= c
FT /bound_molecy= "Probe_99-27943-150"
FT misc_feature 8316
FT /tag= d
FT /note= "Biallelic marker A1"
FT

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FT	primer_bind	/note= "Binds primer 99-27943.pu complement"	FT	primer_bind	/note= "Binds primer 99-24656.tp complement"
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FT	primer_bind	/tag= g	FT	primer_bind	/note= "Binds primer 99-24639.tp"
FT		/note= "Binds primer 99-27935.rp"	FT	primer_bind	160621. .160639
FT		21653. .21671	FT	misc_binding	/tag= af
FT		/tag= h	FT	misc_binding	/note= "Binds primer 99-24639-163. mis"
FT	misc_binding	/note= "Binds primer 99-27935-193. mis"	FT		160628. .160652
FT		21660. .21684	FT	misc_feature	/tag= ag
FT		/tag= i	FT		/bound_moiety= "Probe_99-24639-163"
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FT		21672	FT		/tag= ah
FT		/tag= j	FT	primer_bind	/note= "Biallelic marker A6"
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FT		complement(21673. .21691)	FT		/tag= ai
FT		/tag= k	FT	primer_bind	/note= "Binds primer 99-24639-163. mis complement"
FT	primer_bind	/note= "Binds primer 99-27935-193. mis complement"	FT		160770. .160787
FT		complement(21845. .21864)	FT	primer_bind	/tag= aj
FT		/tag= l	FT	primer_bind	/note= "Binds primer 99-24634.pu"
FT	primer_bind	/note= "Binds primer 99-27935.pu complement"	FT		complement(160785. .160802)
FT		65463. .65471	FT	primer_bind	/tag= ak
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FT		/tag= n	FT	misc_binding	/note= "Binds primer 99-24634-108. mis"
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FT		/tag= p	FT	primer_bind	/note= "Biallelic marker A7"
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FT		complement(65486. .65504)	FT		/tag= ao
FT		/tag= q	FT	primer_bind	/note= "Binds primer 99-24634-108. mis complement"
FT	primer_bind	/note= "Binds primer 8-128-33. mis complement"	FT		complement(161240. .161257)
FT		complement(65856. .65874)	FT	primer_bind	/tag= ap
FT		/tag= r	FT	primer_bind	/note= "Binds primer 99-24634.tp complement"
FT	primer_bind	/note= "Binds primer 8-128.tp complement"	FT		168813. .168830
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FT		/tag= t	FT	misc_binding	/note= "Binds primer 99-7652-162. mis"
FT	misc_binding	/note= "Binds primer 99-31960-363. mis"	FT		168962. .168986
FT		95384. .95408	FT		/tag= as
FT		/tag= u	FT	misc_feature	/bound_moiety= "Probe_99-7652-162"
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FT		95396	FT		/tag= at
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FT		/tag= y	FT		/note= "Binds primer 99-16100.pu"
FT	primer_bind	/note= "Binds primer 99-24656.pu"	FT	primer_bind	170791. .170809
FT		107262. .107280	FT		/tag= ax
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FT		107269. .107293	FT		/tag= ay
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Query Match 61.3%; Score 18.4; DB 5; Length 319608;
* Best Local Similarity 78.6%; Pred. No. 3.8e+02;
Matches 22; Conservative 0; Mismatches 6; Indels 0; Gaps 0;
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Oy 1 CTGACCCCTCTGACCTCGAGTTCG 28
    |||||
gb 8690 CTGACCCATCTCGATTGAGATTACG 8717
```

```
RESULT 13
AAL52198
ID AAL52198 standard; cDNA; 340449 BP.
XX
AC AAL52198;
XX
XX 22-SEP-2003 (first entry)
XX
DE Human secreted protein genomic DNA coding sequence.
XX
XX Human; gene; ds; secreted protein; chromosome 5; tissue typing;
KM secreted protein-related disease; transgenic animal; drug screening;
KM pharmacogenomic analysis; single nucleotide polymorphism; SNP.
XX
XX Homo sapiens.
OS
XX
FH Key location/Qualifiers
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FT      /standard_name= "Single nucleotide polymorphism"
FT      replace(26508, G)
FT      /*tag= ap
FT      /standard_name= "Single nucleotide polymorphism"
FT      replace(26570, A)
FT      /*tag= aq
FT      /standard_name= "Single nucleotide polymorphism"
FT      replace(26582, G)
FT      /*tag= ar
FT      /standard_name= "Single nucleotide polymorphism"
FT      replace(26693, A)
FT      /*tag= as
FT      /standard_name= "Single nucleotide polymorphism"
FT      replace(26884, T)
FT      /*tag= at
FT      /standard_name= "Single nucleotide polymorphism"
FT      replace(27320, T)
FT      /*tag= au
FT      /standard_name= "Single nucleotide polymorphism"
FT      replace(27339, A)
FT      /*tag= av
FT      /standard_name= "Single nucleotide polymorphism"
FT      replace(27542, G)
FT      /*tag= aw
FT      /standard_name= "Single nucleotide polymorphism"
FT      replace(28586, A)
FT      /*tag= ax
FT      /standard_name= "Single nucleotide polymorphism"
FT      replace(28591, T)
FT      /*tag= ay
FT      /standard_name= "Single nucleotide polymorphism"
FT      replace(28599, A)
FT      /*tag= az
FT      /standard_name= "Single nucleotide polymorphism"
FT      replace(30857, A)
FT      /*tag= ba
FT      /standard_name= "Single nucleotide polymorphism"
FT      replace(31503, T)
FT      /*tag= bb
FT      /standard_name= "Single nucleotide polymorphism"
FT      replace(33671, A)
FT      /*tag= bc

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FT      /standard_name= "Single nucleotide polymorphism"
FT      replace(35045, C)
FT      /*tag= bd
FT      /standard_name= "Single nucleotide polymorphism"
FT      replace(35944, G)
FT      /*tag= be
FT      /standard_name= "Single nucleotide polymorphism"
FT      replace(37157, G)
FT      /*tag= bf
FT      /standard_name= "Single nucleotide polymorphism"
FT      replace(37603, T)
FT      /*tag= bg
FT      /standard_name= "Single nucleotide polymorphism"
FT      replace(39242, G)
FT      /*tag= bh
FT      /standard_name= "Single nucleotide polymorphism"
FT      replace(39404, A)
FT      /*tag= bi
FT      /standard_name= "Single nucleotide polymorphism"
FT      replace(40395, C)
FT      /*tag= bj

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Query Match      60.7%; Score 18.2; DB 8; Length 340449;
Best Local Similarity 87.0%; Pred. No. 4.7e+02;
Matches 20; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy      2 TGAACCCCTCTGACTCGAGACT 24
Db      25999 TGAACCCCTCCCACTCCAGAGT 26021

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RESULT 14
ADB74382
ID      ADB74382 standard; DNA; 42325 BP.
XX      AC
XX      ADB74382;
XX      DT
XX      04-DEC-2003 (first entry)
XX      DE
XX      Mycobacterium leprae DNA #16.
XX      KW
XX      Non-naturally occurring peptide; anion pump protein; tuberculosis;
XX      hypersensitivity reaction; tuberculostatic; gene; ds.
XX      OS
XX      Mycobacterium leprae.
XX      PN
XX      US6583266-B1.
XX      PD
XX      24-JUN-2003.
XX      PF
XX      16-SEP-1994; 94US-00311731.
XX      PR
XX      19-AUG-1993; 93US-00109181.
XX      PR
XX      22-OCT-1993; 93US-00142558.
XX      PA
XX      (GENO-) GENOME THERAPEUTICS CORP.
XX      PI
XX      Smith DR, Mao J;
XX      DR
XX      WPI; 2003-656441/62.
XX      PS
XX      Disclosure; SEQ ID NO 131; 26pp; English.
XX      CC
XX      The invention relates to a non-naturally occurring peptide of
XX      Mycobacterium tuberculosis comprising an amino acid sequence
XX      corresponding to an anion pump protein. The invention also relates to a
XX      non-naturally occurring nucleic acid corresponding to a DNA sequence of
XX      Mycobacterium tuberculosis or Mycobacterium leprae. The new peptide is
XX      useful as a vaccine against Mycobacterium tuberculosis or Mycobacterium
XX      leprae or for screening for new tuberculosis drugs. Purified proteins

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CC derived from the sequences of the invention may elicit a specific immune
CC response. The peptide may also be used to detect hypersensitivity
CC reactions of individuals exposed to Mycobacterium tuberculosis or
CC Mycobacterium leprae. The proteins and peptides may be affixed to solid
CC supports to detect antibodies typical of hypersensitivity reactions, from
CC a patient's sera. This sequence represents Mycobacterium leprae DNA of
CC the invention. Note: The sequence data for this patent did not form part
CC of the printed specification but was obtained in electronic format
CC directly from USPTO at seqdata.uspto.gov/sequence.html.

XX
SQ Sequence 42325 BP; 9673 A; 13128 C; 11330 G; 8194 T; 0 U; 0 Other;

Query Match 60.0%; Score 18; DB 10; Length 42325;

Best Local Similarity 80.8%; Pred. No. 5e+02;
Matches 21; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

QY 5 ACCCTCTCGACTCGAGTTCGGCT 30

DB 688 ACCCTCTCGACTCGAGTTCGGCT 713

RESULT 15

ADM27081_10/c

Continuation (11 of 17) of ADM27081 from base 1000001 (Hypertrophile Methanopyrus kan
WP Sequence split into 17 fragments LOCUS ADM27081 Accession Adm27081

Fragment Name	Begin	End
WP ADM27081_00	1	110000
WP ADM27081_01	100001	210000
WP ADM27081_02	200001	310000
WP ADM27081_03	300001	410000
WP ADM27081_04	400001	510000
WP ADM27081_05	500001	610000
WP ADM27081_06	600001	710000
WP ADM27081_07	700001	810000
WP ADM27081_08	800001	910000
WP ADM27081_09	900001	1010000
WP ADM27081_10	1000001	1110000
WP ADM27081_11	1100001	1210000
WP ADM27081_12	1200001	1310000
WP ADM27081_13	1300001	1410000
WP ADM27081_14	1400001	1510000
WP ADM27081_15	1500001	1610000
WP ADM27081_16	1600001	1694968

Query Match 60.0%; Score 18; DB 11; Length 110000;

Best Local Similarity 80.8%; Pred. No. 5.4e+02;
Matches 21; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

QY 1 CTGAGCCCTCTCGACTCGAGATTTC 26

DB 69684 CTTTGCCGCTCTCGACTCGAGATTTC 69659

Search completed: April 26, 2005, 11:11:26
Job time : 433 secs